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Inactivation of histone chaperone HIRA un.masks a link between normal embryonic development of melanoblasts and maintenance of adult melanocyte stem cells

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| <p>Title of the abstract</p> | <p>Inactivation of histone chaperone HIRA unmasks a link between normal embryonic development of melanoblasts and maintenance of adult melanocyte stem cells.</p> |
| <p>Name of authors (presenter <u>underlined</u>) and their affiliations (institution, city, country)</p> | <p><u>Farah Jaber-Hijazi</u>^{1,2,3}, Karthic Swaminathan^{2,4}, Kathryn Gilroy^{1,2}, Alexander T. Wenzel⁵, Anthony Lagnado⁶, Kristina Kirschner^{1,2}, Neil Robertson^{1,2}, Claire Reid^{1,2}, Neil Fullarton^{1,2}, Jeff Pawlikowski^{1,2}, Karen Blyth^{1,2}, Jill P. Mesirov⁵, Taranjit Singh Rai^{1,2,7}, João F. Passos⁶, Laura M. Machesky^{1,2} and Peter D. Adams^{1,2,8}</p> <p>¹ Institute of Cancer Sciences, University of Glasgow, Glasgow, G61 1OH, UK. ² CRUK Beatson Institute, Glasgow, G61 1BD, UK. ³ School of Health and Life Sciences, University of the West of Scotland, Hamilton International Technology Park, Glasgow, G72 0LH, UK ⁴ Centre for Skin Sciences, Faculty of Life Sciences, University of Bradford, Bradford, BD7 1DP, UK. ⁵ Department of Medicine, University of California San Diego (UCSD), La Jolla, CA, USA; Moores Cancer Center, University of California, San Diego, USA, ⁶ Department of Physiology and Biomedical Engineering, Mayo Clinic, Rochester, MN, USA. ⁷ Northern Ireland Centre for Stratified Medicine, Ulster University, Ulster, BT47 6SB, UK. ⁸ Sanford Burnham Prebys Medical Discovery Institute, 10901 North Torrey Pines Road, La Jolla, CA 92037, USA.</p> |
| <p>Abstract (max. 350 words)</p> <p>→ Use font size 8 max.</p> | <p>Histone chaperone Hira, which deposits histone variant H3.3 into chromatin in a DNA replication-independent manner, has been implicated in epigenetic memory and is thought to play a role in both early development and aging. However, there are only sparse data on connections between integrity of embryonic development and healthy aging. The pigmentary system, consisting of differentiated melanocytes and melanocyte stem cells (McSCs) of the adult hair follicle and their precursor melanoblasts in embryos, has been valuable in understanding mechanisms of development, aging and disease. Here, we describe a conditional knockout mouse model, <i>Tyr::Cre Hira^{fl/fl}</i>, in which McSCs, melanocytes and their embryonic melanoblast precursors are specifically deficient for Hira and chromatin deposition of histone H3.3. We find that Hira is required for establishment of normal embryonic melanoblast numbers <i>in vivo</i>, supported by single cell RNA sequencing data, and melanoblast identity <i>in vitro</i>. Despite this, by birth, <i>Tyr::Cre Hira^{fl/fl}</i> mice contain a comparable number of melanocytes as wild type mice, and young adults have normal functioning McSCs and only very mildly hypopigmented hair coat. However, neonate melanoblasts from <i>Tyr::Cre Hira^{fl/fl}</i> mice are sensitive to stress both <i>in vitro</i> and <i>in vivo</i> and exhibit more telomere-associated DNA damage foci, a marker of premature aging, than do those from wild type mice. In line with this, knock out of <i>Hira</i> during embryogenesis in <i>Tyr::Cre Hira^{fl/fl}</i> mice caused a premature defect in adult McSC maintenance and premature hair greying, while inducible knock out of <i>Hira</i> in young adult <i>Tyr::Cre-ERT2 Hira^{fl/fl}</i> mice resulted in no observable defect. These studies of the Hira histone chaperone show that perturbations of <i>in utero</i> embryogenesis can cause only modest phenotypic variations at birth and in young adulthood, but profound abnormalities and features of unhealthy aging in later life.</p> |



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| <p>Presenting Person (name, institute, address and email) → Use font size 8 max.</p> | <p>Dr. Farah Jaber-Hijazi School of Health and Life Sciences, University of the West of Scotland, Hamilton International Technology Park, Glasgow, G72 0LH, UK Farah.jaber@uws.ac.uk</p> |

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